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1 **Structural classification of insecticidal proteins – towards an *in silico* characterization**
2 **of novel toxins.**

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10

11 **Abstract:**

12 The increasing rate of discovery of new toxins with potential for the control of invertebrate
13 pests through next generation sequencing, presents challenges for the identification of the
14 best candidates for further development. A consideration of structural similarities between
15 the different toxins suggest that they may be functionally less diverse than their low sequence
16 similarities might predict. This is encouraging from the prospective of being able to use
17 computational tools to predict toxin targets from their sequences, however more
18 structure/function data are still required to reliably inform such predictions.

19

20 **Introduction:**

21 The insecticidal toxins of *Bacillus thuringiensis*, *Lysinibacillus sphaericus*, *Photorhabdus spp.*
22 and other bacteria represent a rich resource for the control of pest insects. The increasing
23 rate of discovery of new toxins, driven by next-generation sequencing, will expand our arsenal
24 of potential biocontrol agents but this, in itself, presents new challenges. Even with past rates
25 of toxin discovery, toxins have rarely been tested against more than a few species of insects
26 (van Frankenhuyzen, 2009) and, in the future, toxicity testing of large numbers of new toxins

27 against a wide range of insects will not be feasible. To facilitate the selection of toxins for
28 study, different criteria may be applied, including identification of the toxin in a strain known
29 (from a previous screening) to have interesting biocidal activity or relatedness to known
30 toxins. Here we consider the prospects for a further, selective method through the prediction
31 of activity. We highlight some of the challenges that may be encountered and propose steps
32 that will bring us closer to this goal. Useful predictions would not only assist in the selection
33 of toxins for development but would also have value in support of the regulatory process of
34 biopesticide product registration, where the potential to predict off-target activities would be
35 valuable.

36

37 The *B. thuringiensis* nomenclature system (Crickmore et al., 1998) currently contains several
38 hundred individual sequences, divided between 74 classes of Cry toxin, 3 classes of Cyt toxin,
39 4 classes of Vip toxin and one SIP toxin. *L. sphaericus* strains may produce the BinA/B toxin,
40 Mtx1, Mtx2, Mtx3, Mtx4, sphaericolysin, Cry48 and Cry49 (reviewed in (Berry, 2012)) and
41 *Photorhabdus* strains can produce Tc toxins, PirA/B and Mcf toxins (ffrench-Constant et al.,
42 2007). This represents a great diversity of toxins but some simplification can be achieved by
43 considering these proteins in terms of their structural characteristics (known or predicted).
44 Table 1 shows the toxin classes, colour-coded by sequence homology groups. As can be seen,
45 the 3-domain Cry toxins represent the largest structural family (and also encompass the
46 PirA/B toxin, recently shown to be equivalent to a 3-domain toxin with a dissociated
47 domain III (Lee et al., 2015)). There is also a large group of toxins that is rich in beta-sheets
48 with general structural similarity to aerolysin. This group includes Cry46 and toxins
49 identified by Pfam (Bateman et al., 1999) to be members of either the Etx/Mtx2 family or the
50 Toxin_10 family. Other groups include the Cyt toxins, the ADP-ribosyl transferase toxins Mtx1
51 and Vip1/2 (along with the Vip1-like Vip4 protein). Cry34 is an aegerolysin like protein and
52 with Cry35 is part of a two-component toxin (Kelker et al., 2014). Cry37, which itself is part

53 of a two-component toxin with Cry23, which shows structural homology with Cry34 (Rydel et
54 al., 2001). Other toxins, which appear unrelated and have no published structures, are Cry6,
55 Cry22, Cry55, Vip3 and Mcf. Our knowledge of the structure and function of toxins within
56 these groups varies and it will be useful to consider the major groups separately.

57

58 **The 3-domain toxins:** These toxins are the best-characterised, with the first structure
59 published in 1991 (Li et al., 1991) and with several decades of studies on the specificity and
60 mode of action of members of this family. The steps leading to toxicity for this family are well-
61 known and involve ingestion by the invertebrate target, solubilisation of toxin crystals in the
62 gut, proteolytic activation by gut enzymes, one or more receptor binding step, followed by
63 membrane insertion (Pardo-Lopez et al., 2012). Insect specificity could be mediated by any of
64 the above steps, for example changes in proteinase activity (Loseva et al., 2002) but the most
65 important determinants of specificity are the binding to and specificity for receptors on the
66 surfaces of target cells.

67 As suggested by the name of this family, the structure of the active toxin is composed
68 of 3 distinct structural domains. Domain I is formed from a bundle of alpha helices and is
69 involved in pore formation by the toxin. Domain II has a beta prism structure that appears to
70 be related to carbohydrate binding proteins and Domain III has a beta sandwich fold.
71 Domains II and III appear to have roles in receptor binding and specificity of the toxins as
72 demonstrated by domain swapping experiments that have altered target specificity (Lee et al.,
73 1995; Pigott and Ellar, 2007). Bioinformatic analysis suggests that the 3 toxin domains evolve
74 at different rates (Bravo, 1997) and this may have implications for target specificity.

75 Within the 3-domain toxin family, we find toxins with activity against insects in several
76 orders, principally amongst the Lepidoptera with fewer active against the orders Diptera and
77 Coleoptera, and with small numbers active against Hymenoptera and Hemiptera as well as
78 toxins affecting nematodes and gastropods (reviewed recently (Palma et al., 2014a)).

79 Members of this family active against human cancer cells have also been reported (Ohba et al.,
80 2009), although it is clearly unlikely that they have co-evolved with this host. However,
81 correlation between sequence identity and target range is generally poor even when analysis
82 is carried out at the level of the individual domains (de Maagd et al., 2001). This highlights the
83 need for analysis at a level below that of the domains themselves. Within domain II, several
84 exposed loops (the $\alpha 8$ loop, and loops 1, 2 and 3) have been identified as potentially
85 important for receptor binding. The variability of these regions and their dispositions in 3
86 example toxins is shown in Figure 1. These surface loops are amongst the most variable in
87 sequence and in length between individual toxins and even minor modifications have been
88 shown to change targeting (Bravo et al., 2013). For example, Cry4Ba has no significant
89 activity against *Culex* mosquitoes but the substitution of Asp454 in domain II loop 3 with the
90 sequence Pro-Ala-Thr results in high toxicity to *Culex* species without reduction in toxicity
91 towards *Aedes aegypti* (Abdullah et al., 2003). However, these loops may not be the sole
92 domain II mediators of specificity, for example it has been shown that residues remote from
93 these loops (illustrated in Figure 2) contribute to dipteran/lepidopteran specificity in the
94 Cry2Aa/Cry2Ab toxins (Morse et al., 2001).

95 An understanding of the receptor-binding interactions of the 3-domain proteins is
96 made complex by the diversity of putative receptor proteins for these toxins. The most
97 commonly identified binding partners include cadherins, aminopeptidases and alkaline
98 phosphatases (Pigott and Ellar, 2007). However, even for single toxins such as Cry1Ac and
99 Cry4Ba, a large number of interacting proteins can be identified through proteomic studies
100 (Bayyareddy et al., 2009; Krishnamoorthy et al., 2007) (Table 2). Although the significance of
101 these binding interactions for toxicity is not known, we cannot discount possible physiological
102 significance for these interactions, for example prohibitin, identified as a potential Cry4Ba
103 binding-protein has been demonstrated to bind toxin in *Aedes* cells in culture (Kuadkitkan et
104 al., 2012). Glycolipids are also potential receptors for 3-domain toxin binding as shown for

105 Cry1Ac (Garczynski and Adang, 2000), Cry2Ab (Ma et al., 2012) and may be particularly
106 important for the nematode-active Cry5B (Barrows et al., 2007; Griffiths et al., 2005). The
107 diversity of possible receptors and the probable involvement of more than one receptor in
108 toxicity, adds to the challenge of predicting toxin activity.

109 Several studies have attempted to map regions of the toxins that may interact with
110 receptor proteins. For example, residue Tyr445 in loop 3 of Cry1Aa domain II was identified
111 as being important in binding to the cadherin BtR175 of *Bombyx mori* (Atsumi et al., 2005)
112 while Val582 in domain III was shown to be important for interaction with aminopeptidase N
113 (Atsumi et al., 2008). In parallel, there have been studies to map regions of receptors that
114 may interact with the toxins (Pigott and Ellar, 2007) but even when a common class of
115 receptor (eg a cadherin) is considered, there is little correspondence between regions
116 interacting with different toxins.

117

118 **The beta sheet toxins:** There are several classes of toxins in Table 1 that appear to be rich in
119 beta sheets. These include Cry37 (de Maagd et al., 2003) and the aegerolysin-like Cry34
120 (Kelker et al., 2014) which both form part of binary toxins (with Cry23 and Cry35
121 respectively, discussed below). The presence of extended beta sheets also characterises the
122 sphaericolysin/anthrolysin family of toxins that is highly conserved across isolates from
123 *L. sphaericus*, *B. thuringiensis*, *Bacillus cereus* and *Paenibacillus alvei* (Berry, 2012; Bourdeau et
124 al., 2009) and appear to act as cholesterol dependent cytolysins (From et al., 2008; Nishiwaki
125 et al., 2007). At present, the literature lacks sufficient information to allow structure/function
126 predictions for these toxins. However, a number of proteins in Table 1 belonging to the
127 Etx/Mtx2 family, the Toxin_10 family, along with Cry46 are rich in beta sheets and show a
128 general fold similar to aerolysin. This group of toxins is the most numerous after the
129 3-domain toxins and some analysis of structure and specificity is possible. The general
130 structure of these toxins features a head region and a tail region that features long beta

131 strands (available structures for these proteins are shown in Figure 3). Cry45 and Cry46 are
132 produced by *B. thuringiensis* but have no known activity against invertebrates. However,
133 activity against mammalian cancer cells has been demonstrated (Ohba et al., 2009).
134 Structurally, Cry46 most closely resembles the Etx/Mtx2 family (although Pfam analysis of its
135 primary sequence does not assign it to this family). The Etx/Mtx2 family differs from the
136 Toxin_10 family in two clear features. In the Toxin_10 family, the head domains contain beta
137 trefoil motifs similar to carbohydrate-binding domains and these heads are formed
138 exclusively from the N-terminal end of the proteins. The beta trefoil domains may have a role
139 in toxin interactions with glycoproteins or glycolipids to facilitate receptor binding or other
140 stages of the mechanisms of action of the toxins. In contrast, in the Etx/Mtx2 family, the head
141 lacks the beta trefoil and is composed of residues from the N-terminal region and from a
142 further stretch of amino acids much closer to the C-terminus (before the C-terminal sequence
143 completes the last long beta strand of the tail domain). In these features, Toxin_10 proteins
144 resemble toxins such as the haemolytic lectin from the parasitic mushroom *Laetiporus*
145 *sulphureus* (Mancheno et al., 2005), while the Etx/Mtx2 family resembles mammalian toxins
146 such as aerolysin (Figure 3) and epsilon toxin from *Clostridium perfringens*.

147 When the ability of these aerolysin-like toxins to target invertebrate or mammalian
148 cells is compared to their overall sequence, there is no obvious correlation (Figure 4A). When
149 analysis is performed at the level of the phylogeny of individual head and tail domains
150 (Figure 4B and C) there is still no clear correlation, indicating once again the need for deeper,
151 subdomain analysis to be undertaken to predict activity.

152 Understanding the receptor binding and specificity of the Toxin_10 family is further
153 complicated by the existence of partner proteins for many of these toxins. Cry36 is a clear
154 exception since it is reported to act alone to kill *Diabrotica* larvae (Rupar et al., 2000).
155 Another protein in the Toxin_10 family (41.9 kDa protein) may be encoded by *B. thuringiensis*
156 but no toxicity has been discovered to date and no partner protein has been identified (Palma

et al., 2014b). All of the other Toxin₁₀ proteins act with their specific partner proteins to form binary toxins as follows: BinA and BinB (both Toxin₁₀ proteins)(Broadwell et al., 1990; Oei et al., 1990); Cry34 (aegerolysin-like) and Cry35 (Toxin₁₀)(Kelker et al., 2014; Masson et al., 2004); Cry48 (3-domain) and Cry49 (Jones et al., 2007). The role of each protein in these binary pairs is clearly significant to understanding the specificity of the toxins and potential binding of both components presents further challenges to prediction. The binding of both Cry48 and Cry49 to *Culex* brush border membrane fractions has been shown, with Cry49 suggested to be the principal binding component (Guo et al., 2016). In the case of the Bin toxin, BinB appears to be the major receptor-binding component in *Culex* mosquitoes but in *Anopheles* BinA also appears to be able to bind (Charles et al., 1997; Oei et al., 1992). Binding of BinB to target membranes appears to be mediated by residues at its N-terminal end (Oei et al., 1992; Romao et al., 2011; Singkhamanan et al., 2010), consistent with receptor recognition via the head domain (Srisucharitpanit et al., 2014). The BinA/BinB toxin appears to bind to a single toxin receptor, a GPI anchored α -glycosidase (Silva-Filha et al., 1999), which may simplify the investigation of binding interactions (particularly when compared to the complex receptor binding of 3-domain toxins). Both the receptor, Cqm1, from the Bin-sensitive *Culex quinquefasciatus* and the ortholog from the insensitive *Aedes aegypti* are known and a region involved in binding has been identified. This includes a Gly-Gly motif, potentially on a surface loop, which may be required for productive interaction (Ferreira et al., 2014). If binding to single receptors proves to be a general feature of this class of toxin, prediction of activity may be made more straightforward but this may also have implications for the ease with which insects may acquire resistance. A number of the Etx/Mtx2 family of Cry toxins are also reported to require a binary partner for full activity. Cry23Aa (also known as ET33) acts with Cry37 (ET34) and has activity against various coleopteran insects (Donovan et al., 2000; Ekobu et al., 2010). Cry15Aa and Cry33Aa form binary partnerships with two other proteins (40kDa and NT32KD respectively) that, due to their lack of individual activity, have not been

183 assigned Cry names, and share no obvious sequence similarity with any other characterized
184 protein (Kim et al., 2003; Naimov et al., 2011).

185

186 **Convergent evolution of the beta sheet toxins?:**

187 Figure 3 indicates that three of the homology groups highlighted in Table 1 (Toxin_10, Cry46
188 and Etx/Mtx2) share significant sequence similarity and indeed are very similar to toxins such
189 as aerolysin. In the aerolysin family of proteins, the conserved beta sheet dominated
190 “aerolysin fold” is believed to adopt a barrel conformation within a membrane with the
191 associated domains primarily having a binding role (Szczesny et al., 2011). It is conceivable
192 that despite the diversity in primary sequence, many of the non 3-domain Cry toxins could
193 share significant structural/functional homology. This would take the form of the aerolysin
194 tail fold associated with a head domain involved in targeting the protein to a particular
195 receptor. This head domain could be part of the same protein as the aerolysin fold or come
196 from an associated binary partner. A number of Cry and Cyt toxins have acquired a ricin-like
197 beta-trefoil carbohydrate-binding domain and while it is tempting to speculate that this could
198 have given these proteins novel binding activities, there is currently no evidence that this is
199 the case (reviewed by (Adang et al., 2014)).

200

201 **The future for *in silico* analyses:**

202 The ultimate objective of such work is to be able to predict the likely specificity of a toxin from
203 primary sequence data. While the computational power to be able to achieve such a goal is
204 available, the underlying data required to inform such analyses are still lacking. Significant
205 progress has been made in recent years in the elucidation of new toxin structures and the
206 potential to derive reliable structures from modelling approaches is discussed elsewhere in
207 this issue (Berry and Board, 2016). Similarly, while there are a lot of data available on the
208 target specificity of individual toxins (Palma et al., 2014a; van Frankenhuyzen, 2009), many of

209 these are contradictory, thus compounding efforts to derive meaningful associations. The
210 area where least information is known concerns the interaction of the toxin with the target
211 cell (Vachon et al., 2012). As discussed above, even when putative receptors are identified,
212 determining which interactions are crucial for toxicity is far from straightforward. There is a
213 need to elucidate more structures for confirmed receptors, and ideally toxin-receptor
214 complexes, which can then lead to *in silico* predictions of toxin-receptor interactions. A
215 number of studies have used docking analysis to indicate the likely interaction between a Cry
216 toxin and its putative receptor (Ahmad et al., 2015; Tajne et al., 2012; Zhao et al., 2012).
217 Without a detailed understanding of which interactions are crucial for toxicity, such studies
218 are likely to throw up many false positives, indeed one report predicts that Cry1Ac could have
219 activity against cattle (Ebenezer et al., 2013), a prediction that is not supported by
220 experimental observations.

221 In summary, whilst our understanding of toxin structure is rapidly progressing, and might
222 suggest that the large family of toxins is less diverse than was thought, we are still a long way
223 from the goal of being able to match toxins and hosts based on primary sequence data
224 generated from genome sequencing.

225

226

227 Table1: Toxins and their homology groups

228 Toxins of *B. thuringiensis*, *L. sphaericus* and *Photorhabdus* spp. are shown with colouring to

229 indicate homology groups: light blue = 3-domain toxins; peach = Etx/Mtx2 toxins; pink =

230 Toxin_10 family proteins; violet = Cyt toxins; khaki = aegerolysin toxins; grey = ADP ribosyl

231 transferase-related proteins; toxins not falling in to these groups are coloured differently.

Cry1	Cry21	Cry41	Cry61	Vip3
Cry2	Cry22	Cry42	Cry62	Vip4
Cry3	Cry23	Cry43	Cry63	BinA
Cry4	Cry24	Cry44	Cry64	BinB
Cry5	Cry25	Cry45	Cry65	Mtx1
Cry6	Cry26	Cry46	Cry66	Mtx2
Cry7	Cry27	Cry47	Cry67	Mtx3
Cry8	Cry28	Cry48	Cry68	Mtx4
Cry9	Cry29	Cry49	Cry69	Sphaericolysin
Cry10	Cry30	Cry50	Cry70	PirA
Cry11	Cry31	Cry51	Cry71	PirB
Cry12	Cry32	Cry52	Cry72	Mcf
Cry13	Cry33	Cry53	Cry73	
Cry14	Cry34	Cry54	Cry74	
Cry15	Cry35	Cry55	Cyt1	
Cry16	Cry36	Cry56	Cyt2	
Cry17	Cry37	Cry57	Cyt3	
Cry18	Cry38	Cry58	Sip	
Cry19	Cry39	Cry59	Vip1	
Cry20	Cry40	Cry60	Vip2	

232

233 Table 2: Potential receptors for 3-domain toxins.

234 Proteins discovered through proteomics as potential binding partners for two 3-domain

235 toxins, Cry1Ac and Cry4Ba are shown.

Cry4Ba (Bayyareddy et al., 2009)	Cry1Ac (Krishnamoorthy et al., 2007)
Cadherin	Cadherin
Alkaline phosphatases (3)	Alkaline phosphatases
ATPase	ATPases
Actin	Actin
Serine and metallo peptidases	Aminopeptidases
Prohibitin	Desmocollin-like protein
Mitoporin	
Flotillin-1	
ATP synthase	

236

237

238 Figure Legends

239 Figure 1: Domain II loops

240 The structures of Cry1Aa, Cry2Aa and Cry3Aa (PDB codes 1CIY, 1I5P and 1DLC respectively)

241 are shown with domains I and III shown as yellow and brown ribbons while domain III is

242 shown as a surface representation with the exposed regions of the α 8 loop, and loops 1, 2 and

243 3 coloured cyan, orange, green and magenta, respectively.

244

245 Figure 2: Residues implicated in Cry2A specificity

246 The Cry2Aa structure (PDB 1I5P) is shown with domain II in grey. Residues shown to
247 contribute to lepidopteran specificity are coloured blue while those involved in dipteran
248 specificity are shown in magenta.

249

250 Figure 3: Structures of beta sheet toxins

251 The structures of the insecticidal toxins BinB (PDB 3WA1 (Srisucharitpanit et al., 2014)),
252 Cry35 (PDB 4JP0 (Kelker et al., 2014)), Cry23 (PDB 4RHZ), Cry45 (PDB 2D42 (Akiba et al.,
253 2006)), Cry46 (PDB 2ZTB (Akiba et al., 2009)) and Cry51 (PDB 4PKM (Xu et al., 2015)) are
254 shown along with the structures of the haemolytic toxin from *L. sulphureus* (Lsulph: PDB
255 1W3A (Mancheno et al., 2005) and proaerolysin (PDB 1PRE (Parker et al., 1994)). Head
256 regions are coloured cyan, tails blue and the extra domain in proaerolysin in green.

257

258 Figure 4: Phylogenetic relationship of insecticidal and mammalian-active beta sheet toxins

259 The amino acid sequences of the toxins were compared using MEGA6 and phylogenetic trees
260 were built using the maximum likelihood algorithm. For those toxins where the head and tail
261 domains are discontinuous, the separate regions encoding each domain were combined and
262 analysed as a contiguous sequence. Those toxins with known activity against mammals are
263 highlighted in bold red.

264

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 460

Figure 1

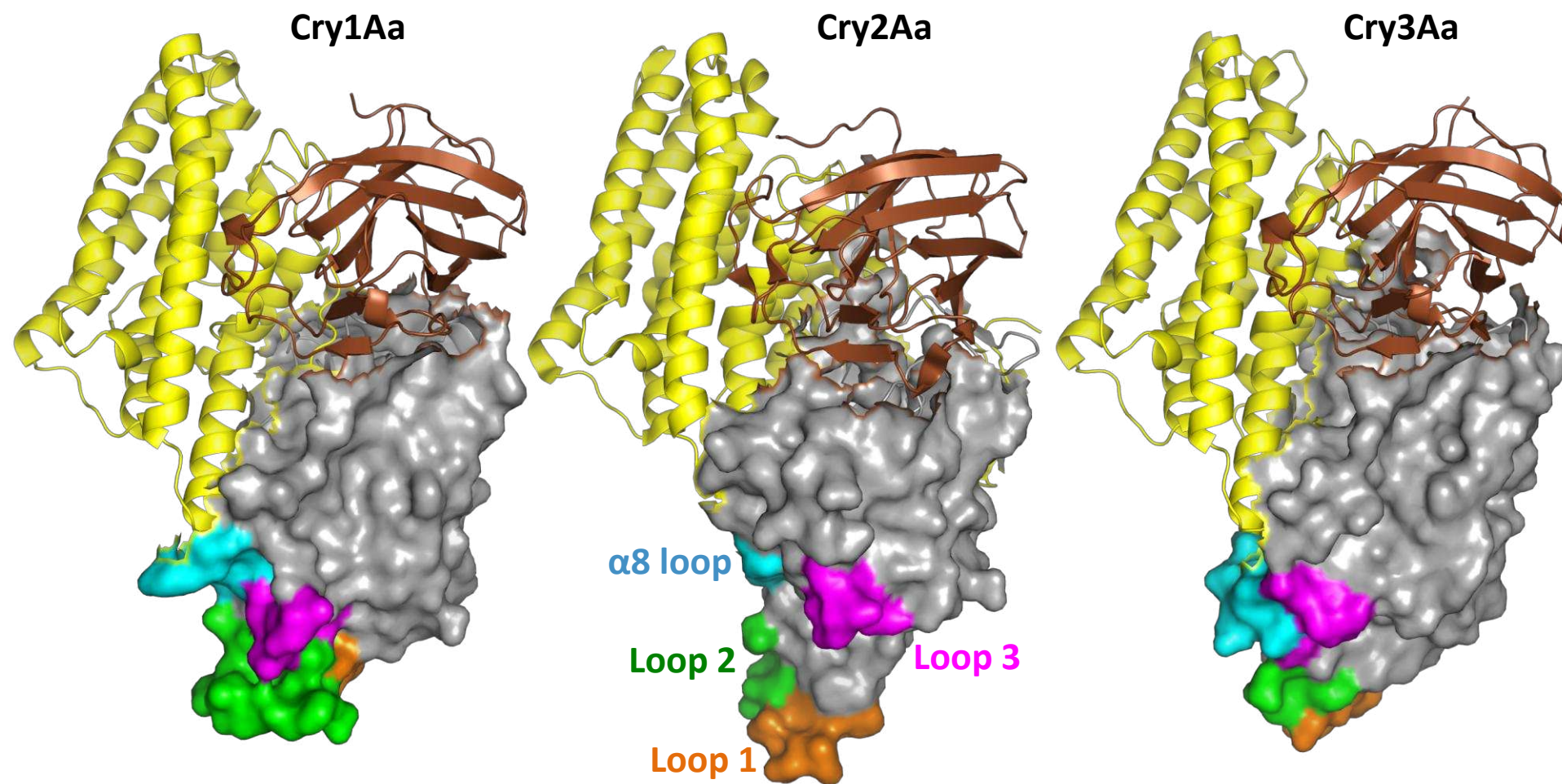


Figure2

Figure 2

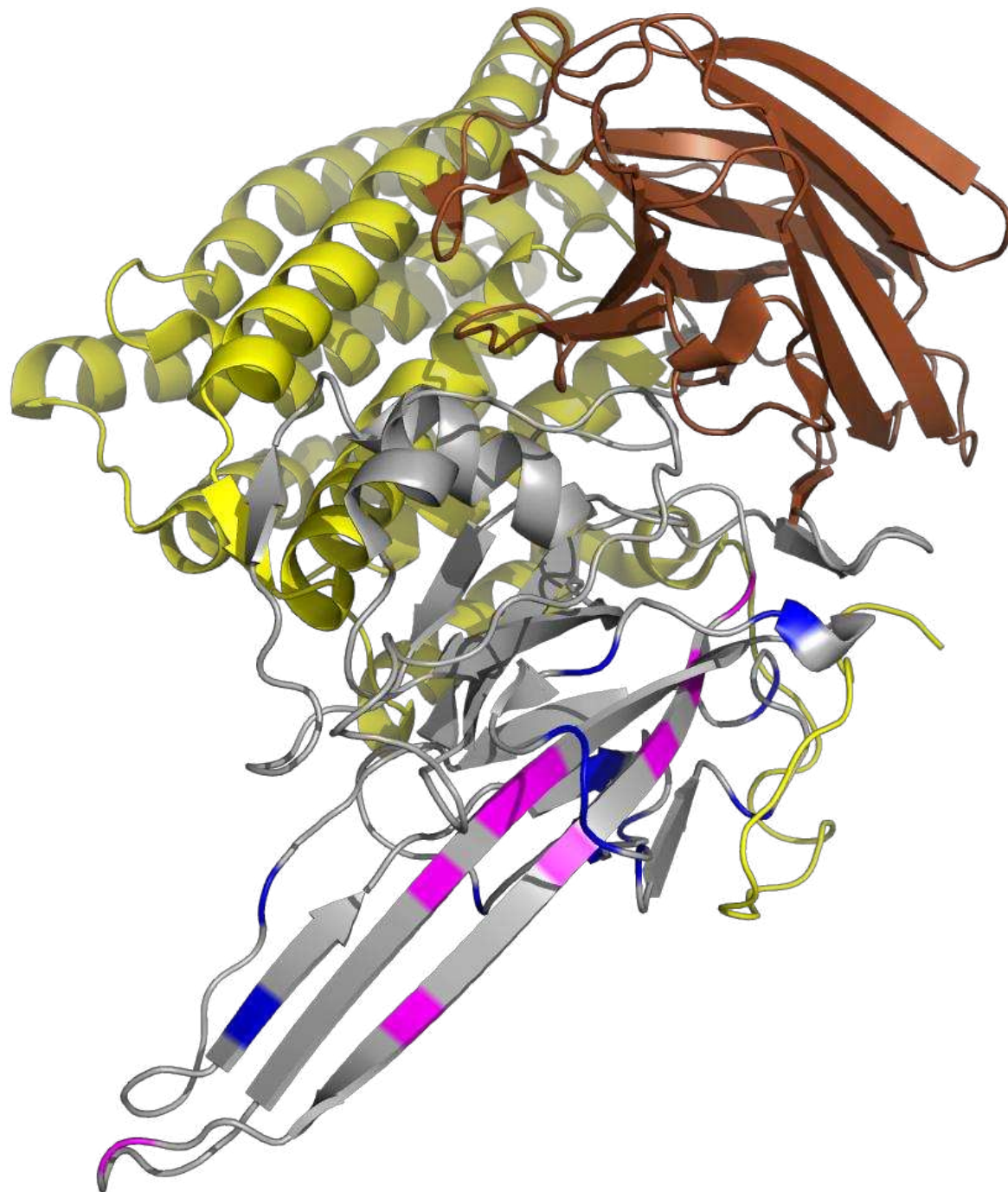


Figure3

Figure 3

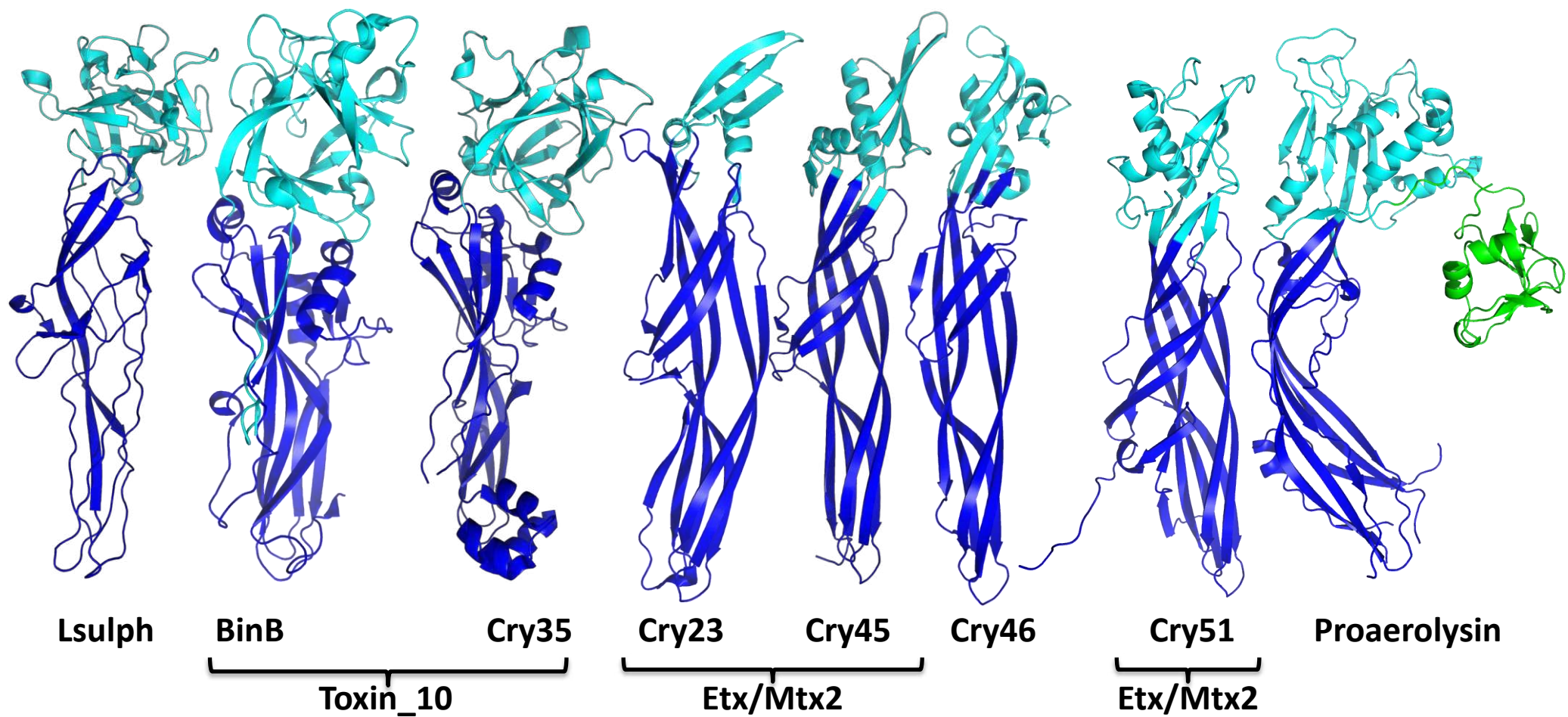
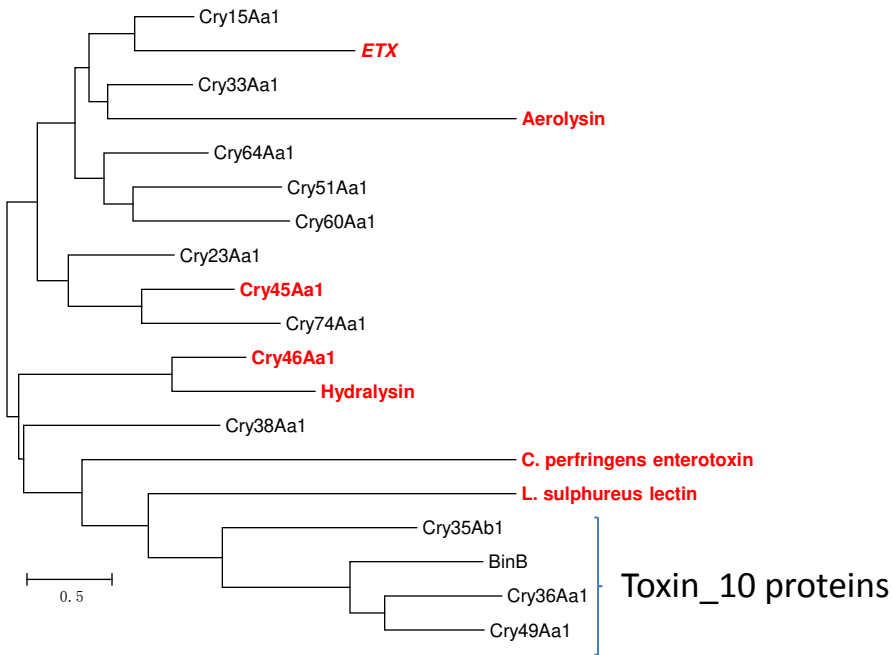


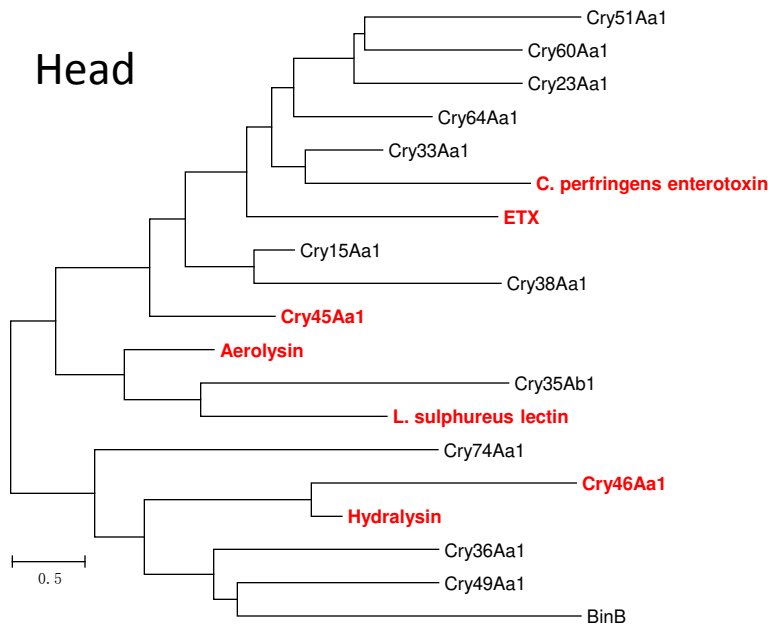
Figure4

Figure 4

A Overall



B Head



C Tail

